

CRITERIA FOR PRIOR AUTHORIZATION**Migraine Acute Treatment Agents**

BILLING CODE TYPE For drug coverage and provider type information, see the [KMAP Reference Codes webpage](#).

MANUAL GUIDELINES Prior authorization will be required for all current and future dose forms available. All medication-specific criteria, including drug-specific indication, age, and dose for each agent is defined in table 1 below.

Celecoxib (Elyxyb)
 Lasmiditan (Reyvow™)
 Rimegepant (Nurtec)
 Ubrogapant (Ubrelyvy)

GENERAL CRITERIA FOR INITIAL PRIOR AUTHORIZATION: (must meet all of the following)

- Must be approved for the indication, age, and not exceed dosing limits listed in Table 1.
- For all agents listed, the preferred PDL drug, which treats the PA indication, is required unless the patient meets the non-preferred PDL PA criteria.
- Medication must be prescribed by or in consultation with a neurologist.
- Patient has a diagnosis of migraines that meet ICHD-3 criteria.¹
- Patient must have <2 migraines per week.^{2,3} If a patient has ≥2 migraines^{2,3} per week or ≥4 migraines per month¹, the patient must be actively on prophylactic treatment.
- Prescriber has determined that the pain intensity of the patient's migraine attacks is moderate or severe.
- Patient must have experienced an inadequate response after a trial (at least 2 weeks) of 2 different triptans at a maximum tolerated dose, OR have a documented intolerance or contraindication to all agents within this class listed in table 2.²⁻⁶
 - Prescriber must provide details of all previous medication trials. Documentation must include the medication name(s), trial date(s) and outcome(s) of the trial (i.e. inadequate response, intolerance or contraindication).
- Patient must have a baseline Migraine Treatment Optimization Questionnaire [mTOQ-5] score of 0.^{6,7}

CRITERIA FOR RENEWAL: (must meet all of the following)

- Dose must not exceed limit in table 1.
- Patient must meet all of the following:
 - Migraines occurring < 2 times per week and < 4 times per month or actively on prophylactic therapy.^{1,3,7}
 - Patient must have an mTOQ-5 score of 5.^{6,7}
- The patient has not required the addition of any other acute treatment agent used for migraine headache.⁶

LENGTH OF APPROVAL (INITIAL AND RENEWAL): 6 months

FOR DRUGS THAT HAVE A CURRENT PA REQUIREMENT, BUT NOT FOR THE NEWLY APPROVED INDICATIONS, FOR OTHER FDA-APPROVED INDICATIONS, AND FOR CHANGES TO AGE REQUIREMENTS NOT LISTED WITHIN THE PA CRITERIA:

- **THE PA REQUEST WILL BE REVIEWED BASED UPON THE FOLLOWING PACKAGE INSERT INFORMATION: INDICATION, AGE, DOSE, AND ANY PRE-REQUISITE TREATMENT REQUIREMENTS FOR THAT INDICATION.**

LENGTH OF APPROVAL (INITIAL AND RENEWAL): 6 months

Table 1. FDA-approved age and dosing limits for Migraine Acute Treatment Agents.⁸⁻¹¹

Agents	Indication(s)	Age	Dosing Limits*
Calcitonin Gene-Related Peptide (CGRP) Receptor Antagonists			
Rimegepant (Nurtec)	Acute treatment of migraine	≥18 years	75 mg orally once in 24 hours.
Ubrogepant (Ubrelvy)	Acute treatment of migraine	≥18 years	200 mg orally per 24 hours.
COX-2 Selective Nonsteroidal Anti-inflammatory Drug (NSAID)			
Celecoxib (Elyxyb)	Acute treatment of migraine	≥18 years	120 mg orally once in 24 hours.
Serotonin 5-HT_{1F} Receptor Agonist			
Lasmiditan (Reyvow™)	Acute treatment of migraine	≥18 years	One dose (50 mg, 100 mg, 200 mg) orally once in 24 hours.

* For a maximum of 4 doses per month.^{1,3,6}

Table 2. Prior Acute Migraine Therapies.^{2,3,6}

Triptans	Combination triptans and NSAIDs
Almotriptan (Axert)	Sumatriptan/naproxen (Treximet)
Eletriptan (Relpax)	
Frovatriptan (Frova)	
Naratriptan (Amerge)	
Rizatriptan (Maxalt)	
Sumatriptan (Imitrex)	
Zolmitriptan (Zomig)	

Notes:

Reyvow	Tablets are available in two strengths: 50 mg, 100 mg. ⁸ The recommended dose of Reyvow is 50 mg, 100 mg, or 200 mg taken orally, as needed. No more than one dose should be taken in 24 hours. A second dose of Reyvow has not been shown to be effective for the same migraine attack. The safety of treating an average of more than 4 migraine attacks in a 30-day period has not been established. ⁸
--------	--

References

1. Headache Classification Committee of the International Headache Society (IHS). The International Classification of Headache Disorders, 3rd edition. Cephalalgia. 2018;38:1-211. Available at <https://ichd-3.org/>.
2. Acute Migraine Headache: Treatment Strategies. Am Fam Physician. 2018 Feb 15; 97(4):243-251. Available at <https://www.aafp.org/afp/2018/0215/p243.html>.
3. Treatment of acute migraine headache. Am Fam Physician. 2011 Oct 1;84(7):738]. Am Fam Physician. 2011;83(3):271-280. Available at <https://www.aafp.org/afp/2011/0201/p271.html>.
4. Headaches in over 12s: diagnosis and management. Updated Feb 2020. Available at: <https://www.nice.org.uk/guidance/cg150>. Accessed 6/5/20.
5. The Acute Treatment of Migraine in Adults: The American Headache Society Evidence Assessment of Migraine Pharmacotherapies. Headache: The Journal of Head and Face Pain 55.1 (2015): 3-20.
6. The American Headache Society position statement on integrating new migraine treatments into clinical practice. Headache: The Journal of Head and Face Pain 59.1 (2019): 1-18. Available at <https://headachejournal.onlinelibrary.wiley.com/doi/epdf/10.1111/head.13456>.
7. Validity and reliability of the migraine-treatment optimization questionnaire. Cephalalgia 29.7 (2009): 751-759.
8. Reyvow (lasmiditan) [prescribing information]. Indianapolis, IN: Lilly USA, LLC; January 2020.
9. Nurtec ODT (rimegepant) [prescribing information]. New Haven, CT: Biohaven Pharmaceuticals Inc; February 2020.

DRAFT PA Criteria

10. Ubrelvy (ubrogepant) [prescribing information]. Madison, NJ: Allergan USA Inc; December 2019.
11. Elyxyb (celecoxib) oral solution [prescribing information]. India: Dr. Reddy's Laboratories Limited; May 2020.

DRUG UTILIZATION REVIEW COMMITTEE CHAIR

PHARMACY PROGRAM MANAGER
DIVISION OF HEALTH CARE FINANCE
KANSAS DEPARTMENT OF HEALTH AND ENVIRONMENT

DATE

DATE